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IPC display formats
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spectra
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applications updated
NEWS 5 MAR 31 LPCI now available as a replacement to LDPCI
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NEWS 14 JUN 06 KOREPAT updated with 41,000 documents
NEWS 15 JUN 13 USPATFULL and USPAT2 updated with 11-character
patent numbers for U.S. applications
NEWS 16 JUN 19 CAS REGISTRY includes selected substances from
web-based collections
NEWS 17 JUN 25 CA/Caplus and USPAT databases updated with IPC
reclassification data
NEWS 18 JUN 30 AEROSPACE enhanced with more than 1 million U.S.
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NEWS 19 JUN 30 EMBASE, EMBAL, and LEMBASE updated with additional
options to display authors and affiliated
organizations
NEWS 20 JUN 30 STN on the Web enhanced with new STN AnaVist
Assistant and BLAST plug-in
NEWS 21 JUN 30 STN AnaVist enhanced with database content from EPFULL
NEWS 22 JUL 28 CA/Caplus patent coverage enhanced
NEWS 23 JUL 28 EPFULL enhanced with additional legal status
information from the epoline Register
NEWS 24 JUL 28 IFICDB, IFIPAT, and IFIUDB reloaded with enhancements
NEWS 25 JUL 28 STN Viewer performance improved
NEWS 26 AUG 01 INPADOCDB and INPAFAMDB coverage enhanced
NEWS 27 AUG 13 CA/Caplus enhanced with printed Chemical Abstracts
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NEWS EXPRESS JUNE 27 08 CURRENT WINDOWS VERSION IS V8.3,

AND CURRENT DISCOVER FILE IS DATED 23 JUNE 2008.

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FILE 'HOME' ENTERED AT 11:17:09 ON 18 AUG 2008

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=> file registry
COST IN U.S. DOLLARS          SINCE FILE      TOTAL
                                ENTRY           SESSION
FULL ESTIMATED COST          0.21            0.21

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FILE 'REGISTRY' ENTERED AT 11:17:24 ON 18 AUG 2008
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DICTIONARY FILE UPDATES: 17 AUG 2008 HIGHEST RN 1041629-70-2

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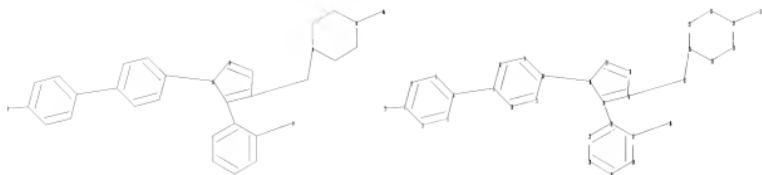
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<http://www.cas.org/support/stngen/stndoc/properties.html>

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Uploading C:\Program Files\STNEXP\Queries\10552065.str



chain nodes :

13 25 32 33

ring nodes :

1 2 3 4 5 6 7 8 9 10 11 12 14 15 16 17 18 19 20 21 22 23 24
26 27 28 29 30 31

chain bonds :

3-13 6-7 10-16 14-32 15-22 23-25 26-32 29-33

ring bonds :

1-2 1-6 2-3 3-4 4-5 5-6 7-8 7-12 8-9 9-10 10-11 11-12 14-15 14-18
15-16 16-17 17-18 19-20 19-24 20-21 21-22 22-23 23-24 26-27 26-31 27-28
28-29 29-30 30-31

exact/norm bonds :

10-16 14-15 14-18 15-16 16-17 17-18 26-27 26-31 26-32 27-28 28-29 29-30
30-31

exact bonds :

3-13 6-7 14-32 15-22 23-25 29-33

normalized bonds :

1-2 1-6 2-3 3-4 4-5 5-6 7-8 7-12 8-9 9-10 10-11 11-12 19-20 19-24
20-21 21-22 22-23 23-24

Match level :

1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:Atom
11:Atom 12:Atom 13:CLASS 14:Atom 15:Atom 16:Atom 17:Atom 18:Atom 19:Atom
20:Atom 21:Atom 22:Atom 23:Atom 24:Atom 25:CLASS 26:Atom 27:Atom 28:Atom
29:Atom 30:Atom 31:Atom 32:CLASS 33:CLASS

L1 STRUCTURE UPLOADED

=> s 11 fam ful
FULL SEARCH INITIATED 11:17:45 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 89 TO ITERATE

100.0% PROCESSED 89 ITERATIONS
SEARCH TIME: 00.00.01

1 ANSWERS

L2 1 SEA FAM FUL L1

=> file caplus
COST IN U.S. DOLLARS

FULL ESTIMATED COST

SINCE FILE TOTAL
ENTRY SESSION
70.11 70.32

FILE 'CAPLUS' ENTERED AT 11:17:51 ON 18 AUG 2008
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FILE COVERS 1907 - 18 Aug 2008 VOL 149 ISS 8
FILE LAST UPDATED: 17 Aug 2008 (20080817/EP)

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=> s 12  
L3           1 L2  
  
=> d 13 ibib abs
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L3 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2008 ACS ON STN
ACCESSION NUMBER: 20041841775 CAPLUS
DOCUMENT NUMBER: 141:350163
TITLE: Preparation of arylpyrazoles as serotonin 5-HT2A and
5-HT2C receptor antagonists
INVENTOR(S): Schiemann, Kai; Ackermann, Karl-August; Arit, Michael;
Finsinger, Dirk; Schadt, Oliver; Van Amsterdam,
Christoph; Bartoszyk, Gerd; Seyfried, Christoph
PATENT ASSIGNEE(S): Merck Patent GmbH, Germany
SOURCE: Ger. Offen., 102 pp.
CODEN: GNXXBX
DOCUMENT TYPE: Patent
LANGUAGE: German
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|--|------|----------|------------------|----------|
| DE 10315572 | A1 | 20041014 | DE 2003-10315572 | 20030405 |
| AU 2004228120 | A1 | 20041021 | AU 2004-228120 | 20040308 |
| CA 2521201 | A1 | 20041021 | CA 2004-2521201 | 20040308 |
| WO 2004089931 | A1 | 20041021 | WO 2004-EP2353 | 20040308 |
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CN, CO, CR, CU, CZ, DE, DK, DM,
DZ, EC, EE, EG, ES, FI, GB, GD, | | | | |

GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC,
 LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI,
 NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY,
 TJ, TM, TN, TR, TT, TZ, UA, US, UZ, VC, VN, YU, ZA, ZM, ZW
 RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ,
 BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE,
 ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI,
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 IE, SI, FI, RO, CY, TR, BG, CZ, EE, HU, PL, SK
 BR 2004009164 A 20060411 BR 2004-9164 20040308
 CN 1768051 A 20060503 CN 2004-80008572 20040308
 JP 2006522035 T 20060928 JP 2006-504584 20040308
 US 20060264419 A1 20061123 US 2005-552065 20051005
 PRIORITY APPLN. INFO.: DE 2003-10315572 A 20030405
 WO 2004-EP2353 W 20040308
 OTHER SOURCE(S): MARPAT 141:350163
 GI

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

AB Preparation of title compds. I [X = CH, N; R1 = H, halo, (CH₂)_nHet, etc.; R2 = (CH₂)_nHet, (CH₂)_nAr, cycloalkyl, etc.; R3, R4 = H, (CH₂)_nCOHET, CHO, etc.; n = 0-5; Ar = (un)substituted Ph; Het = (un)substituted monoarom., bicyclic-heterocycle] and their pharmaceutically acceptable salts were prepared. For example, sodium triacetoxyborohydride mediated reductive amination of 1-methyl-piperazine and aldehyde II, e.g., prepared from 2-fluoro- α , γ -dioxo-benzenebutanoic Et ester in 4-steps, afforded the dihydrochloride salt of arylpyrazole III. In 5-HT2A receptor binding assays, 167-examples of compds. I exhibited IC₅₀ values ranging from 0.015-4.7x10⁻⁷M. Compds. I are claimed suitable as ligands of 5-HT receptors.

| | | | |
|--|------------------|---------------|--|
| => file registry | | | |
| COST IN U.S. DOLLARS | SINCE FILE ENTRY | TOTAL SESSION | |
| FULL ESTIMATED COST | 3.39 | 73.71 | |
| DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) | SINCE FILE ENTRY | TOTAL SESSION | |
| CA SUBSCRIBER PRICE | -0.80 | -0.80 | |

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```
=> s 11 sss ful
FULL SEARCH INITIATED 11:18:30 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED -      163 TO ITERATE

100.0% PROCESSED      163 ITERATIONS          4 ANSWERS
SEARCH TIME: 00.00.01

L4          4 SEA SSS FUL L1

=> file caplus
COST IN U.S. DOLLARS          SINCE FILE      TOTAL
                                ENTRY          SESSION
FULL ESTIMATED COST          178.36         252.07

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) SINCE FILE      TOTAL
                                                ENTRY          SESSION
CA SUBSCRIBER PRICE           0.00          -0.80

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FILE COVERS 1907 - 18 Aug 2008 VOL 149 ISS 8
FILE LAST UPDATED: 17 Aug 2008 (20080817/ED)

Caplus now includes complete International Patent Classification (IPC) reclassification data for the second quarter of 2008.

Effective October 17, 2005, revised CAS Information Use Policies apply. They are available for your review at:

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=> s 14
L5          2 L4

=> d 15 ibib abs 1-2

L5  ANSWER 1 OF 2  CAPLUS  COPYRIGHT 2008 ACS on STN
```

ACCESSION NUMBER: 2004:841775 CAPLUS
 DOCUMENT NUMBER: 141:350163
 TITLE: Preparation of arylpyrazoles as serotonin 5-HT2A and
 5-HT2C receptor antagonists
 INVENTOR(S): Schiemann, Kai; Ackermann, Karl-August; Arlt, Michael;
 Finsinger, Dirk; Schadt, Oliver; Van Amsterdam,
 Christoph; Bartoszyk, Gerd; Seyfried, Christoph
 PATENT ASSIGNEE(S): Merck Patent GmbH, Germany
 SOURCE: Ger. Offen., 102 pp.
 DOCUMENT TYPE: Patent
 LANGUAGE: German
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|--|------|----------|------------------|------------|
| DE 10315572 | A1 | 20041014 | DE 2003-10315572 | 20030405 |
| AU 2004228120 | A1 | 20041021 | AU 2004-228120 | 20040308 |
| CA 2521201 | A1 | 20041021 | CA 2004-2521201 | 20040308 |
| WO 2004089931 | A1 | 20041021 | WO 2004-EP2353 | 20040308 |
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GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC,
LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI,
NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY,
TJ, TM, TN, TR, TT, TZ, UA, US, UZ, VC, VN, YU, ZA, ZM, ZW | | | | |
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ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI,
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| EP 1626967 | A1 | 20060222 | EP 2004-718277 | 20040308 |
| R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
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| CN 1768051 | A | 20060503 | CN 2004-80008572 | 20040308 |
| JP 2006522035 | T | 20060928 | JP 2006-504584 | 20040308 |
| US 20060264419 | A1 | 20061123 | US 2005-552065 | 20051005 |
| PRIORITY APFLN. INFO.: | | | DE 2003-10315572 | A 20030405 |
| | | | WO 2004-EP2353 | W 20040308 |

OTHER SOURCE(S): MARPAT 141:350163
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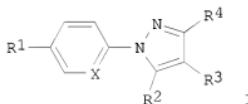
* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

AB Preparation of title compds. I [X = CH, N; R1 = H, halo, (CH2)nHet, etc.; R2 = (CH2)nHet, (CH2)nAr, cycloalkyl, etc.; R3, R4 = H, (CH2)nCOHet, CHO, etc.; n = 0-5; Ar = (un)substituted Ph; Het = (un)substituted monoarom., bicyclic-heterocycle] and their pharmaceutically acceptable salts were prepared. For example, sodium triacetoxyborohydride mediated reductive amination of 1-methyl-piperazine and aldehyde II, e.g., prepared from 2-fluoro- α , γ -dioxo-benzenebutanoyl Et ester in 4-steps, afforded the dihydrochloride salt of arylpyrazole III. In 5-HT2A receptor binding assays, 167-examples of compds. I exhibited IC50 values ranging from 0.015-4.7x10⁻⁷M. Compds. I are claimed suitable as ligands of 5-HT receptors.

ACCESSION NUMBER: 2004:841772 CAPLUS
 DOCUMENT NUMBER: 141:332186
 TITLE: Preparation of arylpyrazoles as serotonin 5-HT2A
 and/or 5-HT2C receptor antagonists.
 INVENTOR(S): Schadt, Oliver; Arlt, Michael; Finsinger, Dirk;
 Schiemann, Kai; Van Amsterdam, Christoph; Bartoszyk,
 Gerd; Seyfried, Christoph
 PATENT ASSIGNEE(S): Merck Patent GmbH, Germany
 SOURCE: Ger. Offen., 78 pp.
 DOCUMENT TYPE: Patent
 LANGUAGE: German
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|--|------|----------|------------------|------------|
| DE 10315569 | A1 | 20041014 | DE 2003-10315569 | 20030405 |
| AU 2004228124 | A1 | 20041021 | AU 2004-228124 | 20040310 |
| CA 2521227 | A1 | 20041021 | CA 2004-2521227 | 20040310 |
| WO 2004089932 | A1 | 20041021 | WO 2004-EP2453 | 20040310 |
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GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC,
LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI,
NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY,
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| EP 1611122 | A1 | 20060104 | EP 2004-718926 | 20040310 |
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| CN 1768052 | A | 20060503 | CN 2004-80008603 | 20040310 |
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| AT 364601 | T | 20070715 | AT 2004-718926 | 20040310 |
| ES 2287710 | T3 | 20071216 | ES 2004-718926 | 20040310 |
| US 2007010531 | A1 | 20070111 | US 2005-552064 | 20051005 |
| PRIORITY APPLN. INFO.: | | | DE 2003-10315569 | A 20030405 |
| | | | WO 2004-EP2453 | W 20040310 |

OTHER SOURCE(S): MARPAT 141:332186
 GI



AB Title compds. [I; R1 = H, A, halo, $(\text{CH}_2)_n\text{Ar}$, cycloalkyl, CF3, NO2, cyano, C(NH)NOH, OCF3; R2 = $(\text{CH}_2)_n\text{Het}$, $(\text{CH}_2)_n\text{Ar}$, cycloalkyl, CF3; R3, R4 = H, $(\text{CH}_2)_n\text{CO}_2\text{R}$, $(\text{CH}_2)_n\text{COH}\text{et}$, CHO, $(\text{CH}_2)_n\text{OR}$, $(\text{CH}_2)_n\text{Het}$, CH:NOA, etc.; R5 = H, A; A = alkyl, alkoxy, alkenyl, alkoxyalkyl; Ar = (substituted) Ph; Het = (aromatic) mono- or bicyclic heterocyclic, heteroatom-containing organic residue; X

= N, CH; with provisos], were prepared. Thus, [1-(4'-fluorobiphen-4-yl)-5-furan-2-yl-1H-pyrazol-4-ylmethyl]methyl(1-methylpyrrolidin-3-yl)amine showed 5-HT2A activity with IC₅₀ = 5.14E-10.

=>